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Sixth Quarterly Report



IC FABRICATION USING ELECTRON-BEAM TECHNOLOGY

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Period Covered

1 December 1977 - 1 March 1978

Contract No. DAAB07-76-C-8105



Technical Support Activity
U.S. Army Electronics Research and Development Command
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This project has been accomplished as part of the U.S. Army Manufacturing and Technology Program, which has as its objective the timely establishment of manufacturing processes, techniques or equipment to ensure the efficient production of current or future defense programs.

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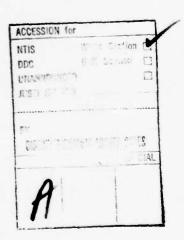
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SECTION I PURPOSE

The overall objective of the program is to implement e-beam writing technology for the fabrication of microcircuits. The technical and economic impact of electron-beam direct slice printing will be demonstrated on 256-bit bipolar RAMs. The elimination of mask masters, masks, and the masking process will eliminate the most significant source of yield loss. This will permit greater circuit design complexity and flexibility which will lead to lower device costs with increased reliability. The complete implementation program is divided into three tasks. Task A, Yield Improvement Through Direct E-Beam Writing, is directed toward developing the manufacturing technology required for e-beam writing with existing equipment and existing resist processes and demonstrating the yield benefits of this technique. Task B, Cost Reduction for E-Beam Writing Through High Speed Resist Implementation, is directed toward implementing identified high speed e-beam resists in order to significantly decrease cycle time and thus reduce the IC bar cost. Task C, Cost Reduction for E-Beam Writing Through Automatic Beam Diameter Control and Automatic Handling, is directed toward utilizing EBMIII's capability of computer-controlled beam size (large and small) on high density circuit (≤0.1 mil) geometries. This program also included implementation of an automated handling system for slices to reduce cycle time and thus further reduce bar cost.

SECTION II RESULTS

A. INTRODUCTION

During this quarter, several basic decisions and changes in direction were made along with some very significant accomplishments. Lot 3 of 74S301A 256-bit bipolar RAMs was dc tested and none of the units passed all the tests. Most units passed the continuity tests but none of them passed the functional tests. The units were retested with the continuity and functional tests removed so that the device parameters could be measured. All of the tested parameters were excellent. An examination of this data, along with a vigorous visual examination of the slices, indicated the failures were caused by defects in the e-beam resist, specifically the positive resist PBS. This also correlates with pinhole data taken using PBS which indicates an abnormally high number of defects when using as an oxide etch barrier. For this reason, a T1 proprietary positive electron resist (T1-313), whose development was proceeding in parallel to the contract, was phased in during this quarter. The pinhole data on this resist (after oxide etch) is comparable with that measured on the best negative photoresists such as Kodak's 747. This T1-313 resist has also allowed plasma etching at each oxide removal step in the process. In addition, a change was made from the double-level metal 256-bit bipolar RAM (54S300) to the single-level metal 256-bit bipolar RAM (74S301A). This change was made because the Tl-Houston production facility was achieving extremely low yields on the 54\$200/300 and had discontinued production. In addition to starting e-beam lots of the 74S301A 256-bit bipolar RAM, photomask lots were also started on the same device. This will allow a yield comparison between standard parts and e-beam delineated devices processed in the same facility. In addition, Dr. Jack Reynolds replaced Mr. Ron Williamson as chief investigator on this contract during this quarter.

B. PROCESS DEVELOPMENT

In a previous quarterly report¹, it was demonstrated that a major yield loss problem for e-beam fabrication of the 54S200/300 256-bit bipolar RAM was the high defect density generated when using PBS electron resist as an oxide etch mask. This quarter a great deal of emphasis has been placed on implementing another positive electron resist (Tl-313) and its processing. Tl-313 was chosen because of its speed, its compatibility with plasma processing and its low defect density. In addition, a plasma etching process compatible with Tl-313 has been worked out for oxide etching at DUF and isolation.

^{1.} IC Fabrication Using Electron-Beam Technology, Contract No. DAAB07-76-C-8105, Fifth Quarterly Report (Dallas, Texas: Texas Instruments Incorporated).

A second major problem that caused difficulty with the early lots of material which were processed was the wash out of etched alignment markers during epi growth. Attempts to realign and pattern another marker set by e-beam led to significant gain and offset errors between the DUF level and the remaining levels. An alternative scheme has been developed and tested successfully which uses contact printing from two photomasks generated on EBMII to place markers on the slices before and after epi growth.

I. TI-313 Resist

Significant progress was made this quarter in implementing TI-313 positive e-beam resist and its processing to replace PBS resist for patterning the 74S301A 256-bit bipolar RAM. The details of manomer synthesis and polymerization have been well established and a high molecular weight (100K MW), narrow dispersivity, copolymer is being provided for formulation into a resist (TI-313). This resist has been evaluated for coating thickness, pinhole density, sensitivity, resolution, etch resistance, adhesion, thermal flow and thermal stability.

Xylene is used as a solvent in the formulation of Tl-313 to provide the optimum combination of solubility, wettability, evaporation rate and viscosity that produces uniform defect-free coatings by spinning. A solids content in the range of 10-14% is convenient for film thicknesses in the range of 0.3-1.2 μ m. A spin speed versus thickness chart for a 10% solution of Tl-313 is shown in Figure 1.

Because of the etch defect problems with PBS, evaluation of TI-313 for pinhole density was one of the first tests undertaken. The procedure for visual counting of etched defects that was described in the previous quarterly report was used. For film thicknesses of $1.0 \,\mu m$ or greater, pinhole densities of $2/cm^2$ or less were found. This compares quite favorably with standard photoresists and much better than PBS as was demonstrated last quarter.

Exposure tests on TI-313 indicate that the critical dose for proper linesize is in the range of 2 to $5 \,\mu\text{C/cm}^2$ depending on thickness and development. Figure 2 shows a plot of the size of a nominally 200 μ inch window as a function of dose for two different development times. Doses much in excess of $6 \,\mu\text{C/cm}^2$ lead to significant amounts of crosslinked material which is not removed by development. This causes some problems in areas that are multiply scanned such as over alignment markers or areas where patterns overlap slightly because of pattern generator errors.

Patterns were developed in TI-313 by spraying for 90 seconds with a 511 mixture of 2-ethoxyethanol/2.6-dimethyl-4-heptanone followed by a 15-second rinse with 2-propanol. One of the disadvantages of TI-313 appears to be that the ratio of developing rates of unexposed to exposed resist is very low. Thus, while clearing out exposed patterns down to the substrate, 50% of

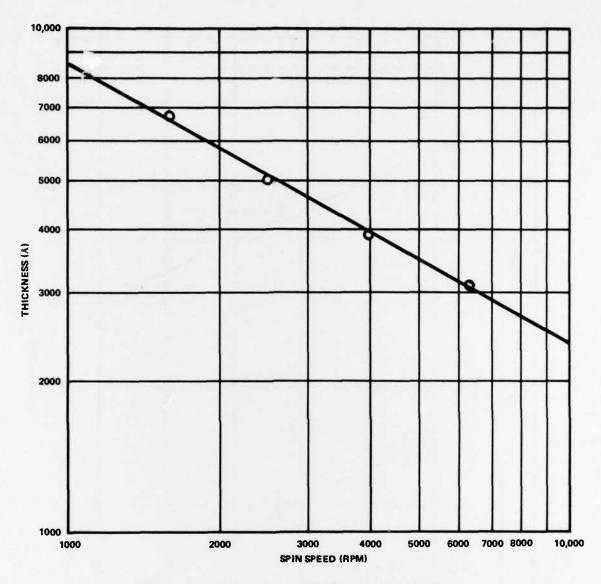


Figure 1. Spin Speed versus Thickness - TI-313 10%

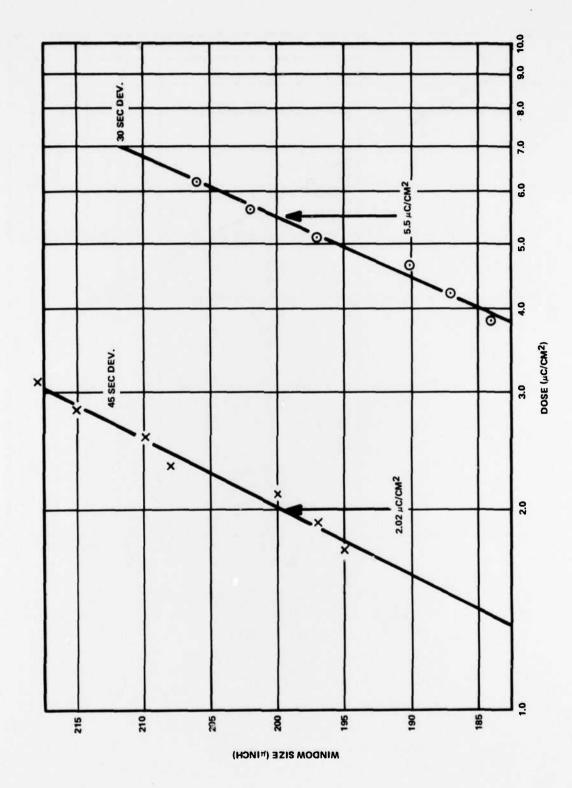


Figure 2. TI-313 Resist Developed Image Size versus Dosc

the film thickness in the unexposed areas is lost. This can be a serious disadvantage in applications where step coverage, pinhole density and removal rates during etching are important.

Thermal analysis of TI-313 films have shown that the material crosslinks in the region of 170°C and begins thermal decomposition in air at about 190°C. Also patterns in the resist begin to flow at 160°C. The baking of the resist after coating and after development is done in a three-zone, belt oven at temperatures of 100-120-140°C in each zone respectively for 15 minutes.

The use of the process described above for TI-313 has been used successfully for delincating the pattern geometries required for fabrication of the 74S301.

2. Plasma Etching

Despite the useful pattern generation capability of TI-313, its adhesion to SiO² is poor and severe undercutting occurs when using TI-313 as a wet etch mask in buffered HF. On the other hand, TI-313 has excellent stability in plasma etching and its removal rates (50Å/min shielded, tubular type, 2000Å/min unshielded parallel plate type) are much lower than PBS or PMMA and are comparable with some negative photoresists.

A process has been developed to use TI-313 at the DUF and isolation levels to mask CF_4/O_2 etching in a shielded tubular reactor. The SiO_2 is etched down to about 1500-2000Å in the plasma and the remaining oxide is etched in buffered HF. This process yields patterns with excellent edge profiles. A brief outline of the process to be used with starting slices of 6000Å of oxide is given below:

- 1. Backside oxide strip
- 2. Coat TI-313, 2500 RPM
- 3. Bake, 100-120-140°C, 15 min
- 4. Repeat steps 2 and 3
- 5. Expose pattern
- 6. Spray develop, rinse, dry
- 7. Inspect and measure
- 8. Bake, 100-120-140°C, 15 min
- 9. Plasma descum, O₂, 1.5 torr, 100 W, 2 min, no tunnel

- 10. Plasma etch, CF₄/O₂ (4%), 1.0 torr, 300 W, 24 min, in tunnel
- 11. Etch to clear, buffered HF, approx. 2 min
- 12. Inspect
- 13. Plasma strip, O₂, 1.5 torr, 300 W, 15 min, no tunnel

Patterns of the DUF and isolation levels etched by this process are shown in Figures 3 and 4. Figure 5 is an SEM micrograph showing the edge profile produced by this etching.

The CF_4/O_2 plasma etching process described above cannot be used at the base, emitter or contact levels because there are at least two different thicknesses of oxide to etch at each level. When the thinnest oxide has completed etching, the silicon beneath is exposed to the very vigorous CF_4/O_2 plasma and is etched at ten times the rate of the remaining oxide. Work is currently underway to solve this problem by using a C_4F_8 plasma etch in a parallel plate reactor of the Reinberg² design. In some cases, conditions can be found where SiO_2 will etch at 4-5 times the rate of Si.



Figure 3. DUF Level (74S301A)

^{2.} U.S. Patent, 3,757,733 issued to Texas Instruments Incorporated.

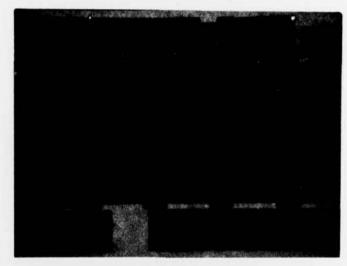
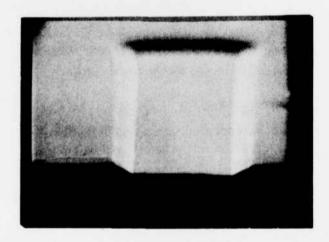


Figure 4. Isolation Level (74S301A)



SEM OF COMBINATION PLASMA ETCH/WET ETCH PROCESS WITH TI-313

Figure 5. 256-Bit Bipolar RAM-DUF Level – 9000Å Oxide

3. Alignment Markers and Masks

Previous reports have demonstrated the wash out of alignment marks during epi growth and the difficulties encountered in trying to do automatic e-beam alignments of the post-epi levels to the pre-epi DUF level. A new method was developed which uses two photomasks fabricated on EBMIII to optically print markers which are etched into the slice. The first mask is used for pre-epi markers to which the DUF level is aligned and the second mask is optically aligned after epi to the DUF level to give markers for alignment of all subsequent levels. Results from the use of this scheme on the first lots of 74S201A to be fabricated have shown realignment accuracies of $\pm 1 \mu m$ between the DUF isolation levels. This is more than adequate for DUF alignment since this is normally the least critical tolerance.

Besides the first two alignment mark masks which are used as described previously, a third mask has been generated which can be used to clear oxide residues which remain in the vicinity of the e-beam markers after etching. These residues are a result of e-beam resist which is crosslinked during the alignment scans and not removed by development. Modification to the alignment scan routine and modifications to the TI-313 electron resist are being investigated as alternate solutions to the problem.

C. SLICE PROCESSING

1. General Discussion

All effort on building the 54S200/S300 at the TI production facility in Houston has ceased due to extremely low yields caused by a circuit design problem. In view of this fact, the implementation of e-beam writing technology with the 250-bit bipolar RAM has been redirected to 74S201A/S301A. This single-level metal 256-bit bipolar RAM is currently being built in Houston with a sufficiently high yield to make it suitable for the e-beam program.

In order to properly evaluate the e-beam processing system for building the 74S201A/S301A, each lot of material processed will be accompanied by a lot processed by conventional contact print resist techniques. Indeed several lots of contact print lots will be processed in the SREL pilot line facility in order to establish a yield capability data point.

2. Process Description

The process to be used for building the e-beam version of the 74S201A/S301A is given in detail in Table I and Figure 6. The contact print monitor lots are processed exactly on the e-beam material except for the resist steps which will be done by conventional contact printing. The e-beam resist steps listed in Table I are discussed in detail in Section 11.B.2.

Table I. Process Flow for E-beam 74S201A/74S301A

Step

Substrate

Type:

Dopant:

Boron

Resistivity:

10-20 ohm-cm

Orientation:

1-1-1 3"

Diemeter:

1st Marker Mask

Piranhe Clean

Spin Swab/Inspect

Bake:

30 minutes

Blue M oven

200° C

Coat:

3.5K RPM

Waycoet III

23 cps for 7KÅ

Softbake: N₂ Dry:

60 minutes 30 minutes plus Vacuum oven

80°C

A&E:

Tower 114

B.0 seconds

Develop/Rinse/

Dry:

Stoddard/ButyI/N2

Inspect:

Silicon Etch

SREL E-Beem Leb

3 **First Oxidation**

Pirenha Clean

Spin Sweb/Inspect

Oxide Temp:

1300° C cycled from 850° C

Time:

360 minutes O₂ 4 I/minutes

Gas: Furnace:

Silicon Tube/Boat

Inspect

Measure Oxide

Thickness:

6400A

Backside Strip OR

Coat:

5.0K RPM

Waycoat III IMS Oven

23 cps for 6400A 65°C

Softbake: A&E:

10 minutes Blank Mask

8.0 seconds

Develop

Inspect Hardbake:

10 minutes

Etch Backside:

7.5 minutes Cold DI H2O IMS Oven Sell 2 5 minutes 125°C 30° C 3X

Rinse: Spray Rinse/

Dry:

5/4 minutes

Inspect Piranha Clean

DUF OR SREL E-Beem Lab

Table 1. Process Flow for E-beam 74S201A/74S301A (Continued)

Step

DUF Diffusion

Pirenhe Cleen and edd "P" type pilots

Inspect

Deglaze:

30 seconds

10% HF

Room Tempereture

Spin Arsenic:

2 minutes 165°C

(Will remove approximetely 170A)

Linear Circuits

Bake:

Inspect Deposition

Temp:

1100°C cycled from 850°C

Time: Gas:

50 minutes

02N2

O₂ 150 cc/min N₂ 2.85 1/min

DUF Window:

Field Oxide:

1050A

7450Å

Inspect

Strip Pilot:

Steam Deglaze:

49% HF

1 minute

Measure and Record

Resistivity Spec:

23-27 Ω/sq.

Temp:

1000°C cycled from 850°C

Time:

15-30-5

Gas:

O2-Steam-N2

O₂ 2 1/min N₂ 2 l/min

DUF Window:

4265A

Field Oxide:

A000A

Strip Pilot and Read

Rs Spec: Deglaze:

29-35 Ω/sq.

4.0 minutes

10% HF

Room Temperature

Avg. Etch Rate

Per Minute:

DUF Window: Field Oxide:

418Å

450 A

Oxide Left for

Drive:

Drive:

DUF Window:

2500A

Field Oxide:

6200A

Temp:

1300°C cycled from 850°C

Time:

300 minutes

Gas:

O₂ 2 1/min

DUF Window:

7000A

Field Oxide:

8300A

Strip Pilot:

Oxide After Drive:

Read Rs and Xj Spec:

Rs 12-17 Ω/sq.

49% HF

35-45 HG Lines Xi 3.0 minutes

Strip Slices

Groove one slice to check if eny penetretion end/or damage.

Epi

Pirenha Cleen

Spin Sweb

Epi:

HCI Etch:

4 HG Lines

Thickness:

.10 - .11 mils .28 - .32 Ω/cm

Resistivity: Evaluate and Record R&T

Table I. Process Flow for E-beam 74S201A/74S301A (Continued)

Stap

2nd Marker Mask

Repeat Step 2, except use 2nd marker mask

2nd Oxidation

Pirenhe Cleen

Spin Sweb/Inspect

Oxide:

Temp:

Time:

1000°C cycled from 850°C

15-70-5

Ges:

O₂-Steam-N₂

02-2 I/min N2-2 I/min

Inspect

Measure Oxide

Thickness:

5400A

Backsida Strip OR 10

Repeat Step 4

Isolation OR SREL E-Baam Lab 11

3rd Marker Mask OR

Piranha Clean

Inspect

Bake: Coat:

30 minutes

5K RPM

10 minutes

Tower 114

Blue M oven

200°C

Waycoet III IMS Oven

23 cps for 6000A 65° C

8.0 seconds

A&E: Develop

Softbake:

Inspect

Hardbake:

10 minutes

To cleer 2nd marker windows

IMS Oven

125°C

Etch: Inspect

Piranhe

Isolation Diffusion 13

Inspect

Deposition:

Temp: Source:

1100°C cycled from 850°C

BBr₃ 25-45-5

750°C

Time: Gas:

N2O2 - N2O2N2 - N2O2

O2-200 cc/min

N₂ Source-30 cc/min

N2-7 1/min

Steem Deglaze:

Temp:

Time: Gas:

10-20-5

O2-Steem-N2

02-2 I/min N2-2 I/min

Degleze:

2.0 minutes

10% HF

30°C

Meesure Rs

on Pilot:

4-5 Ω/sq.

Table I. Process Flow for E-beam 74S201A/74S301A (Continued)

Step						
13	Isolation Diffusion (Continued)					
	Drive:	Temp:	1100°C cycled from 850°C			
		Time:	30-10-5			
		Gas:	O ₂ -Steem-N ₂	O ₂ -2 I/min N ₂ -2 I/min		
	Inspect					
	Meesure Oxide					
	Thickness:	3200-3400A				
	Strip Pilot end					
	Read Rs	5-7 Ω/sq.				
	Xj Pilot:	10-11 HG Lines	· · · · · · · · · · · · · · · · · · ·			
14	Backside Strip OR					
	Repeet Step 4					
15	Base OR SREL E-Beam Lab					
16	3rd Marker Mask OR					
	Repeat Step 12					
17	Base Diffusion					
	Inspect:	Add "N" Pilots				
	Pre-Heat:	Temp:	700°C			
		Source:	Boron Nitride			
		Time:	10 minutes			
		Gas:	N ₂ 2 I/min			
	Deposition:	Temp:	950°C			
		Source:	Boron Nitride			
		Time:	45 minutes			
		Ges:	N ₂ 2 I/min			
	Steam Degleze:	Temp:	750°C			
		Time:	10-20-5			
		Gas:	O ₂ -Steam-N ₂	O ₂ -2 I/min N ₂ -2 I/min		
	Deglaze:	2.0 minutes	10% HF 30°C			
	Measure Rs					
	on Pilot:	$58-66 \Omega/\text{sq}$.				
	Inspect	•	105000 - 1-1 (05000			
	Drive:	Temp: Time:	1050° C cycled from 850° C 30-15-75			
		Gas:		0 01/:-		
		Gas:	O ₂ -Steem-N ₂	O ₂ -2 I/min N ₂ -2 I/min		
	Inspect					
	Measure Oxide					
	Thickness:	3200-3400A				
	A 1 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4					

170-190 Ω/sq .

4.5-5.5 HG Lines

Strip Pilot and

Read Rs: Xj Pilot:

Table I. Process Flow for E-beam 74S201A/74S301A (Continued)

Step

18 **Backside Strip OR** Repeat Step 4 Emitter OR SREL E-Beam Lab 19 3rd Marker Mask OR Repeet Step 12 21 **Emitter Pilot Deposition** Inspect Deposition: Temp: 1000°C Source: POCI3 Time: 5-9-1-4-1 Gas: $N_2O_2 - N_2O_2N_2 - N_2O_2 - N_2O_2 + Steam \\ N_2O_2$ O2-300 cc/min N₂ Source-300 cc/min N₂-2 1/min Measure Oxide Thickness: 2000Å Strip Pilot and Read Rs: 5-7 Ω/sq. Xj Pilot: 3-4 HG Lines Backside Strip OR Repeet Step 4 Contact OR SREL E-Beam Lab 23 24 Piranha Emitter Pilot Anneal/Probe 25 450°C Anneal: Temp: 5-60-5 Time: 4.5 l/min } Gas: Argon-H2-Argon Argon O₂ Scales H_2 Probe: NPN BE 30-60 **Emitter Lot Deposition** Repeet Steps 21 thru 24 Evaporation Piranha Clean Inspect Pletinum Sputter: 500A inspect Aque Regia: 10 minutes inspect Prienha Clean

Table I. Process Flow for E-beam 74S201A/74S301A (Concluded)

Step				
27	Evaporation (Continued)			
	Spin Swab/Inspect			
	TiW Sputter:	1500Å		
	Aluminum Evap:	Thickness:	55 microinches	
		Substrate Temp:	200°C	
	Measure Aluminum			
	Thickness			
	Inspect			
28	Aluminum/TiW Removal SI	REL E-Beem Lab		
29	J-100 Clean			
30	Sinter/Probe			
	Sinter:	Temp:	450°C	
		Time:	60 minutes	
		Gas:	0 ₂ N ₂	O ₂ 50 cc/min N ₂ 2 I/min
	Probe:	NPN βF	30-60	
31	Nitride Deposition			
	Spray Rinse/Spin			
	Dry:	Cold DI H ₂ O		
	Inspect			
	Deposition:	Temp:	230°C	
		Thickness:	3000A	
32	Nitride OR SREL E-Beem L	.eb		
33	Etch and Ash			
	Etch:	8.0 minutes } 100°C		
	Ash:	25 minutes		
34	J-100 Clean			
35	Multi Probe: CDD Measure	ements Lab		

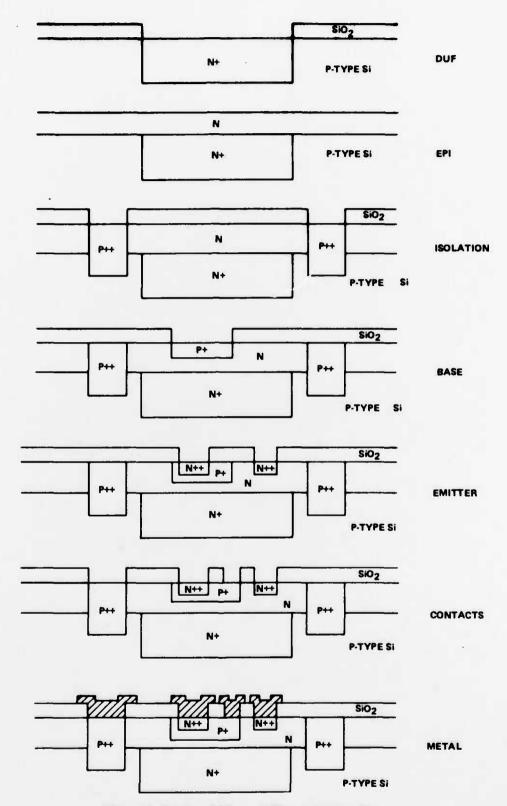


Figure 6. Sectional View of Slice at Various Steps

SECTION III MANPOWER

The following professionals worked on this program 1 December 1977 to 1 March 1978. The percentage of time worked is also shown.

Mr. P. L. Whelan	20%
Mr. R. A. Williamson	50%
Dr. G. L. Varnell	10%
Dr. J. L. Bartelt	50%
Dr. R. A. Owens	50%
Dr. J. Reynolds	50%
Dr. R. A. Robbins	Consultant
Mr. C. D. Winborn	Consultant

In addition, three technicians worked on the program.

